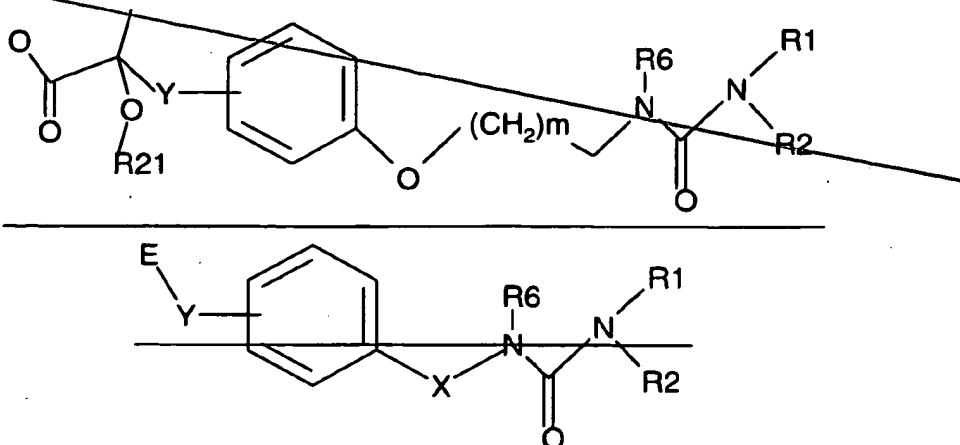


AMENDMENTS TO THE CLAIMS

1. (currently amended) Compound of the structural formula I:

Formula I



and pharmaceutically acceptable salts, solvates and hydrates thereof, wherein:

- (a) R<sub>1</sub>, R<sub>2</sub> and R<sub>6</sub> are each independently selected from the group consisting of hydrogen, C<sub>1</sub>-C<sub>8</sub> alkyl, substituted C<sub>1</sub>-C<sub>8</sub> alkyl, aryl-C<sub>0-4</sub>-alkyl, substituted aryl-C<sub>0-4</sub>-alkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, substituted C<sub>3</sub>-C<sub>6</sub> cycloalkyl, heteroaryl-C<sub>0-4</sub>-alkyl, substituted heteroaryl-C<sub>0-4</sub>-alkyl, C<sub>3</sub>-C<sub>6</sub> cycloheteroalkylaryl-C<sub>0-2</sub>-alkyl, substituted C<sub>3</sub>-C<sub>6</sub> cycloheteroalkylaryl-C<sub>0-2</sub>-alkyl, C<sub>3</sub>-C<sub>6</sub> cycloalkylaryl-C<sub>0-2</sub>-alkyl and substituted C<sub>3</sub>-C<sub>6</sub> cycloalkylaryl-C<sub>0-2</sub>-alkyl; wherein the substituents for said substituted alkyl, arylalkyl, cycloalkyl, heteroarylalkyl, cycloheteroalkylarylalkyl, and cycloalkylarylalkyl are from one to three substituents each independently selected from R<sub>1</sub>';
- (b) R<sub>1</sub>', R<sub>3</sub>', and R<sub>4</sub>' and R<sub>19</sub>' are each independently selected from the group consisting of H, C<sub>1</sub>-C<sub>5</sub> alkyl, C<sub>1</sub>-C<sub>5</sub> alkoxy, C<sub>1</sub>-C<sub>5</sub> haloalkyl, C<sub>1</sub>-C<sub>5</sub> haloalkoxy, nitro, cyano, CHO, hydroxyl, aryl-C<sub>0-5</sub>-alkoxy, aryl-C<sub>0-5</sub>-alkyl, alkylcarboxamido and COOH;
- ~~(c) X is an optionally substituted C<sub>1</sub>-C<sub>5</sub> alkylene linker wherein one carbon atom of the linker may be replaced with O, NH or S;~~
- (d) Y is C, O, S, NH or a single bond; and
- (d) m is 0, 1, or 2; and

(e) R21 is selected from the group consisting of phenyl, substituted phenyl, and C<sub>1</sub>-C<sub>6</sub> alkyl.

- (e) E is selected from the group consisting of hydrogen, C(R3)(R4)A, A, and (CH<sub>2</sub>)<sub>n</sub>-COOR19; wherein said (CH<sub>2</sub>)<sub>n</sub>-COOR19 is optionally substituted with a group selected from C<sub>1</sub>-C<sub>5</sub> alkyl, arylC<sub>0</sub>-C<sub>5</sub>alkoxy, and arylC<sub>0</sub>-C<sub>5</sub>alkyl; and wherein

- ~~(i) n is 0, 1, 2 or 3,~~
- ~~(ii) A is selected from the group consisting of carboxyl, C<sub>1</sub>-C<sub>3</sub>alkylnitrile, carboxamide, sulfonamide, substituted sulfonamide, acylsulfonamide, substituted acylsulfonamide, tetrazole and substituted tetrazole;~~
- ~~(iii) R3 is selected from the group consisting of H, C<sub>1</sub>-C<sub>5</sub> alkyl, and C<sub>1</sub>-C<sub>5</sub> alkoxy, wherein said alkyl and alkoxy are each optionally substituted with from one to three substituents each independently selected from R3';~~
- ~~(iv) R4 is selected from the group consisting of H, halo, C<sub>1</sub>-C<sub>5</sub> alkyl, C<sub>1</sub>-C<sub>5</sub> alkoxy, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, aryl C<sub>0</sub>-C<sub>4</sub> alkyl, and arylC<sub>0</sub>-C<sub>2</sub>alkoxy, or R3 and R4 are optionally combined to form a C<sub>3</sub>-C<sub>4</sub> cycloalkyl, and wherein said alkyl, alkoxy, cycloalkyl, arylalkyl, and arylalkoxy are each optionally substituted with from one to three substituents each independently selected from R4'; and~~
- ~~(f) R19 is selected from the group consisting of hydrogen, arylmethyl, and C1-C4alkyl, wherein said arylmethyl and C1-C4alkyl, are each optionally substituted with from one to three substituents each independently selected from R19'.~~

2. (canceled).
3. (canceled).
4. (canceled).
- <sup>2</sup> 5. (currently amended) A compound as claimed by ~~any one of~~ Claims 1, 2, 3 or 4 wherein R6 is selected from the group consisting of hydrogen, substituted C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> alkyl, substituted aryl-C<sub>0-4</sub>-alkyl, and aryl-C<sub>0-4</sub>-alkyl.
6. (canceled).
7. (canceled).
8. (canceled).
- <sup>2</sup> 8. (currently amended) A compound as claimed by ~~any one of~~ Claims 1, 2, 3, 4, 5, 6, or 7 wherein Y is C.
10. (canceled).
- <sup>4</sup> 11. (currently amended) A compound as claimed by ~~any one of~~ Claims 1, 3, 5, 8, or Claim ~~9~~<sup>3</sup> or 10 wherein aryl is substituted phenyl.
- <sup>5</sup> 12. (currently amended) A compound as claimed by ~~any one of~~ Claims 1 through 11 ~~11~~<sup>4</sup> wherein R2 is hydrogen and R1 is substituted phenyl.
- <sup>6</sup> 13. (original) A compound as claimed by Claim 1 or ~~12~~<sup>5</sup> wherein substituted phenyl is substituted with a group selected from aryl, aryloxy, and arylalkyloxy.
14. (canceled).
15. (canceled).
- <sup>7</sup> 16. (currently amended) A compound as claimed by ~~any one of~~ Claims ~~1 through 15~~<sup>4</sup> wherein the ~~E-CO<sub>2</sub>C((CH<sub>3</sub>)(OR21))~~-Y group is in the para position in relation to the X linker.
- <sup>8</sup> 17. (currently amended) A compound as claimed by ~~any one of~~ Claims 1 through 10, Claim 14, Claim 15, or Claim ~~16~~<sup>7</sup> wherein R1 is selected from unsubstituted phenyl and substituted phenyl, and R6 is hydrogen.
- <sup>4</sup> 18. (currently amended) A compound as claimed by ~~any one of~~ Claims 1 through 17 ~~17~~<sup>7</sup> wherein R1 is substituted phenyl wherein the phenyl substituent is one or two independently selected from the group consisting of CF<sub>3</sub>, C<sub>1</sub>-C<sub>4</sub> alkyl, and halo.
- <sup>10</sup> 19. (Currently amended) A compound as claimed by ~~any one of~~ Claims 1 through 18 Claim ~~18~~<sup>9</sup> wherein R1 is substituted phenyl and R2 is hydrogen.
20. (canceled).

11 ~~21~~. (currently amended) A pharmaceutical composition, comprising a pharmaceutically acceptable carrier and at least one compound as claimed by ~~any one of~~ Claims 1-20.

12 ~~22~~. (currently amended) A method of modulating a peroxisome proliferator activated receptor, comprising the step of contacting the receptor with at least one compound as claimed by ~~any one of~~ Claims 1-20.

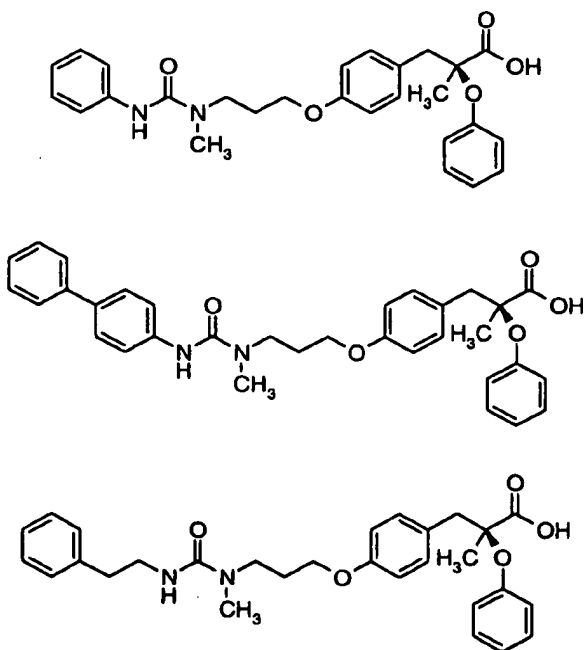
13 ~~23~~. (currently amended) A method of treating diabetes mellitus in a mammal, comprising the step of administering to the mammal in need thereof, a therapeutically effective amount of at least one compound of Claims 1-20.

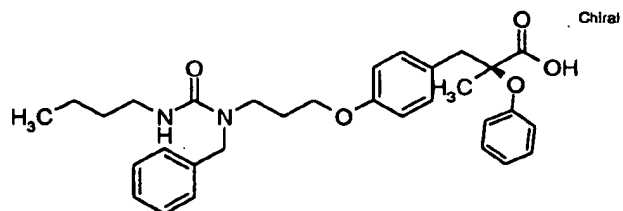
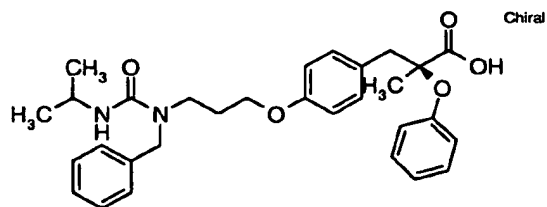
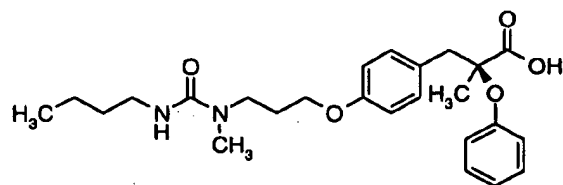
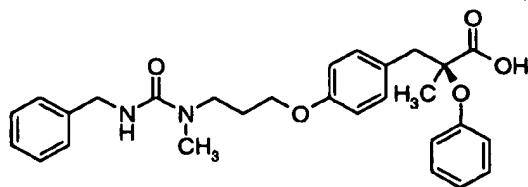
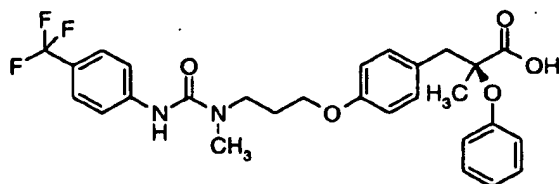
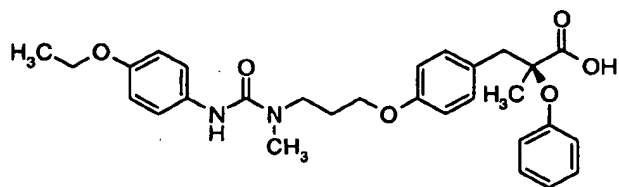
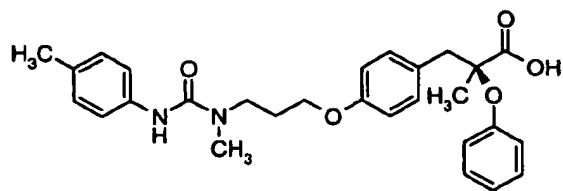
14 ~~24~~. (currently amended) A method of preventing diabetes mellitus in a mammal, comprising the step of administering to the mammal in need thereof, an effective amount of at least one compound of Claims 1-20.

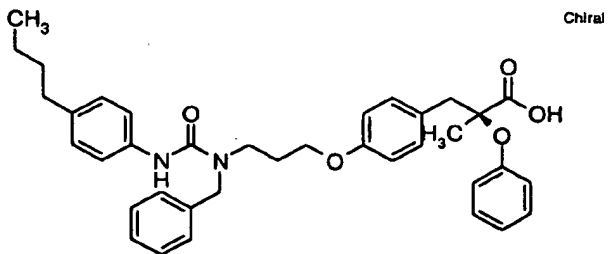
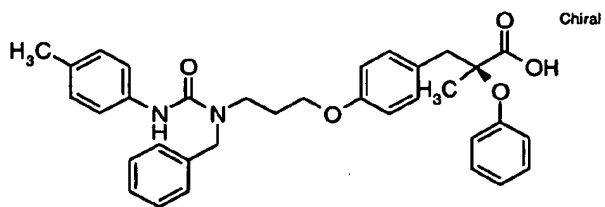
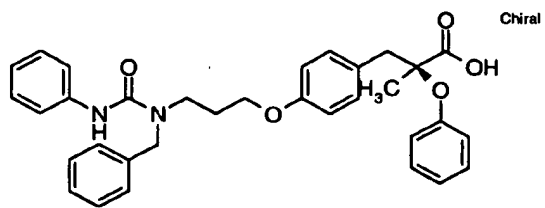
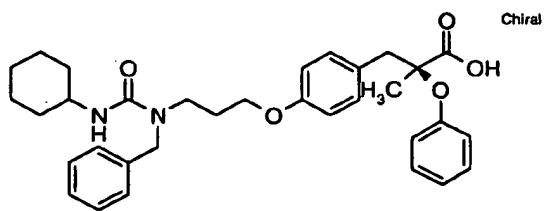
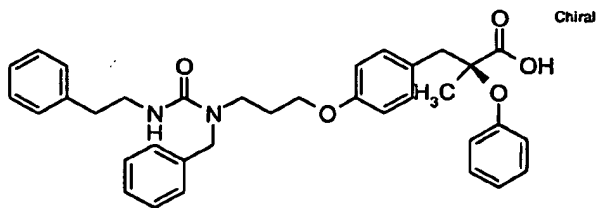
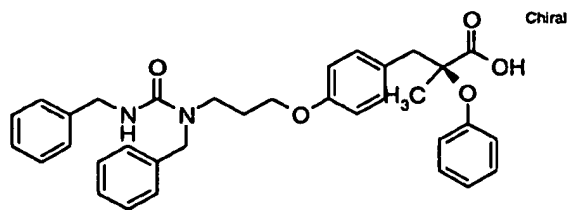
15 ~~25~~. (currently amended) A method of treating Syndrome X in a mammal, comprising the step of administering to the mammal in need thereof, a therapeutically effective amount of at least one compound of Claims 1-20.

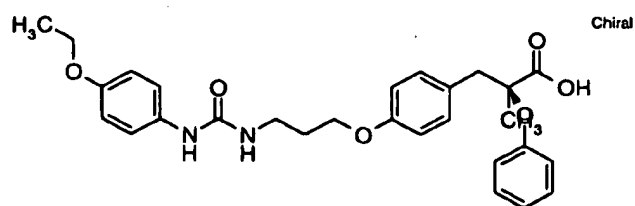
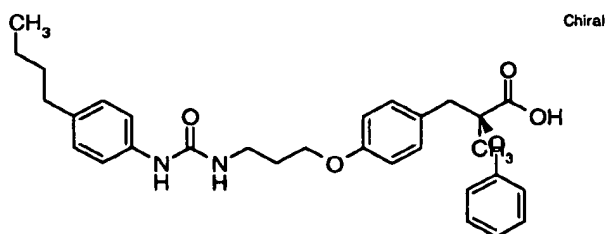
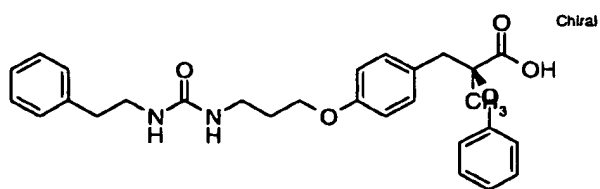
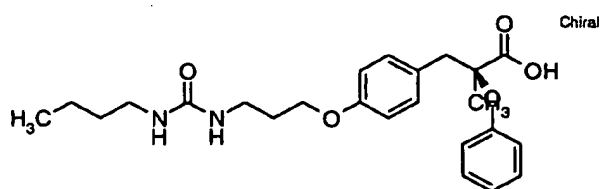
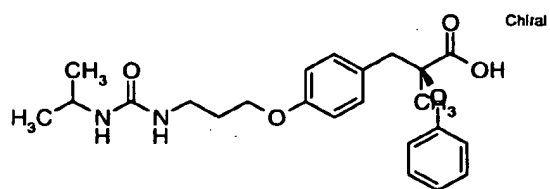
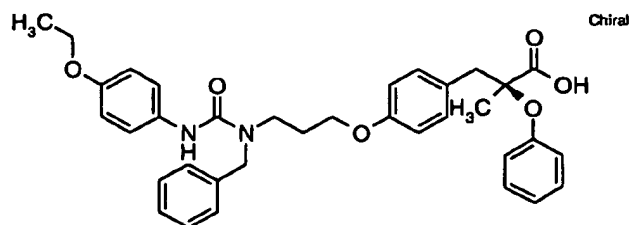
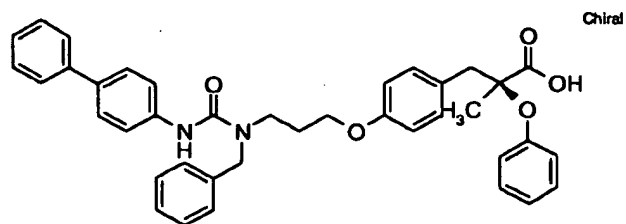
26. (canceled).

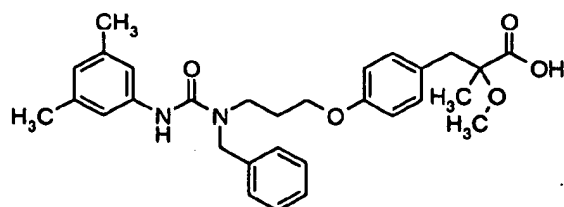
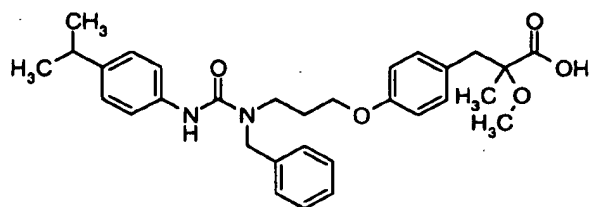
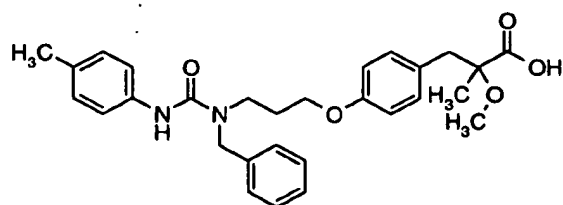
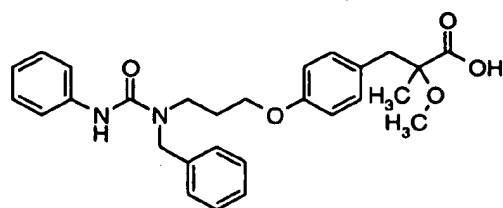
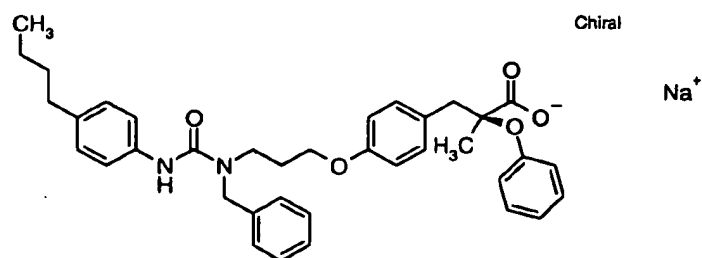
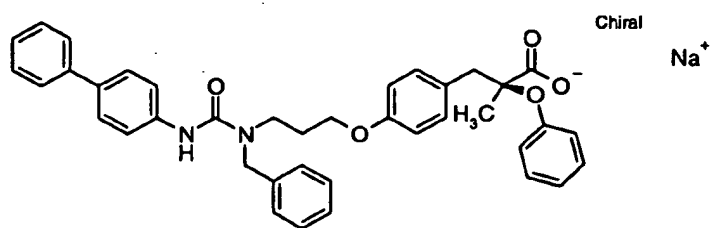
16 ~~27~~. A compound of Claim 1 ~~as disclosed by any one of the examples herein.~~ selected from the group consisting of:



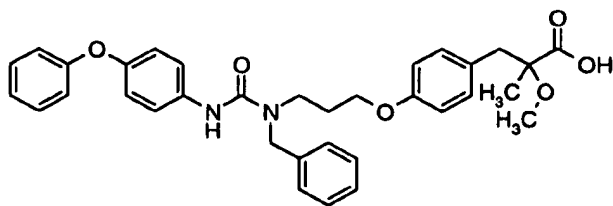
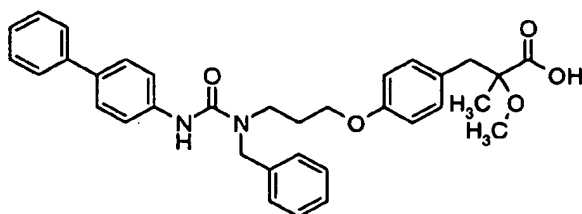
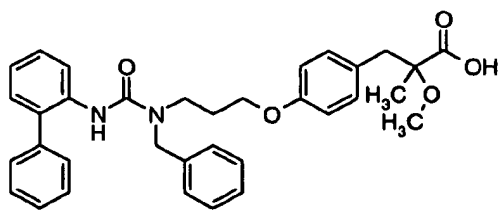
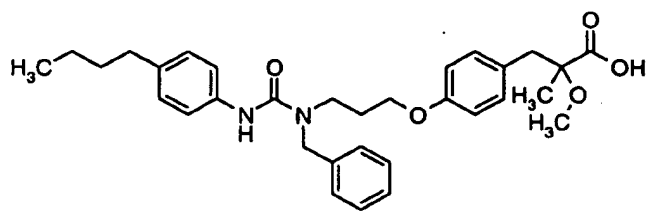
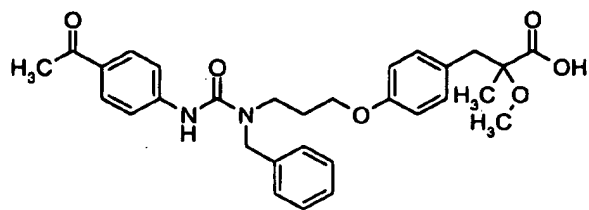
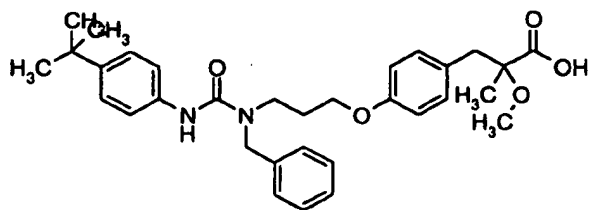


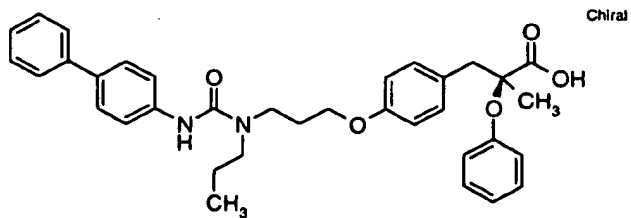
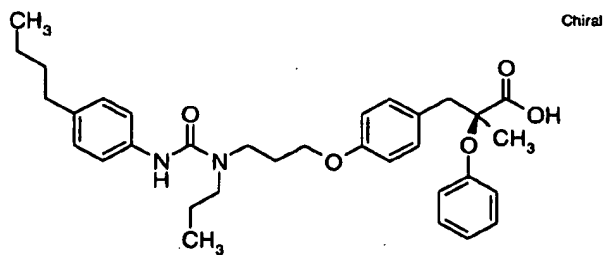
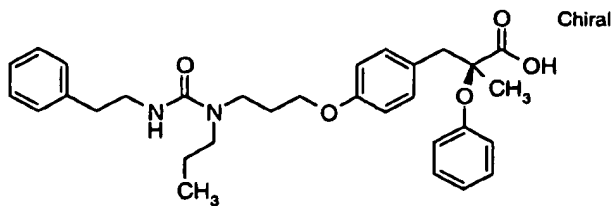
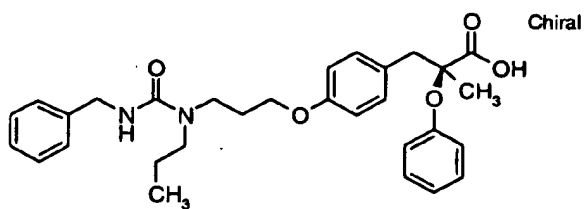
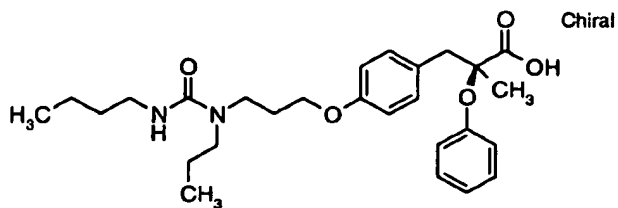
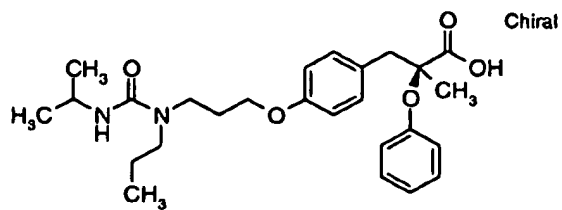


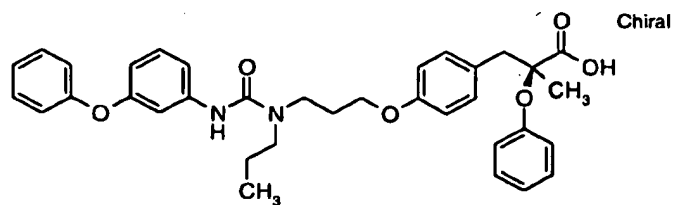
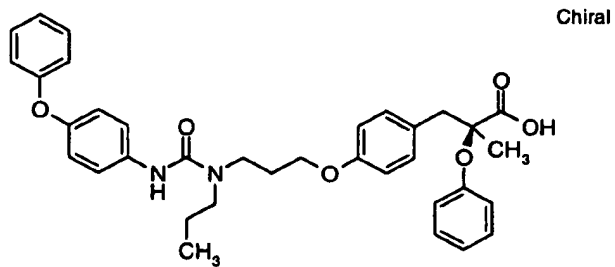
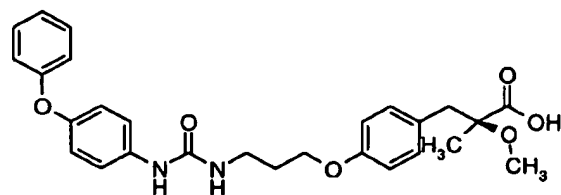
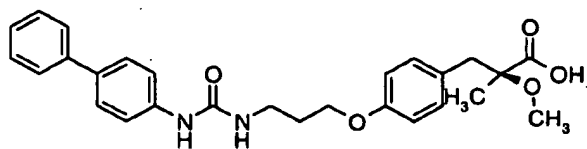
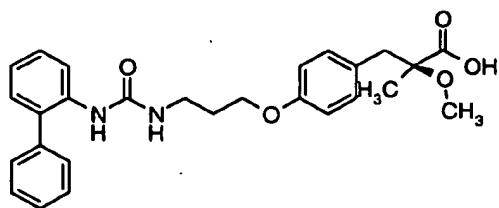
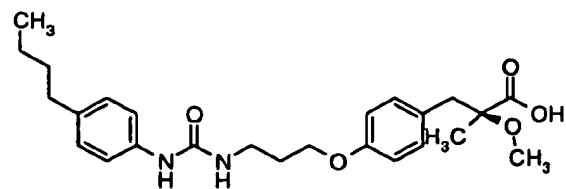
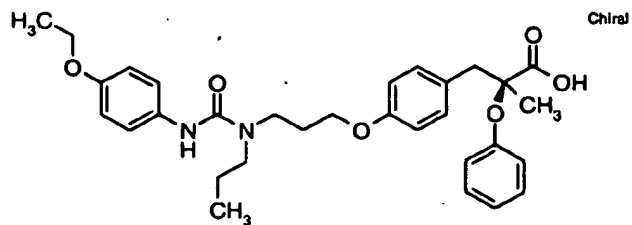


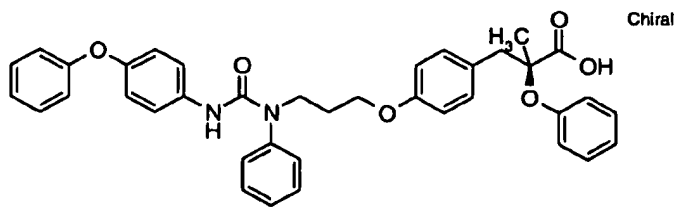
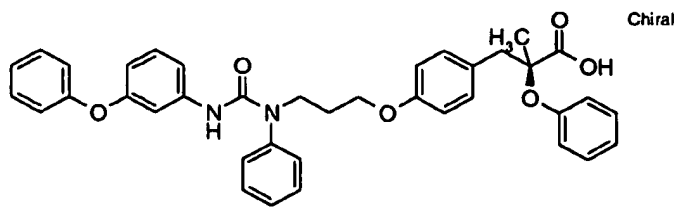
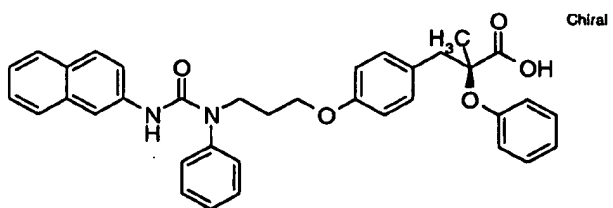
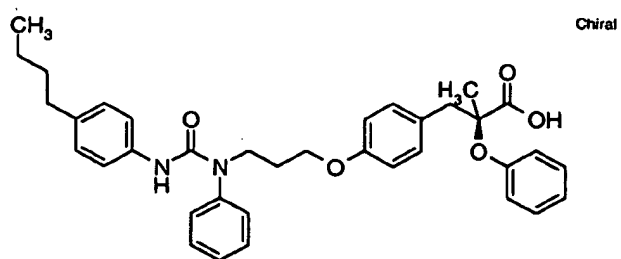
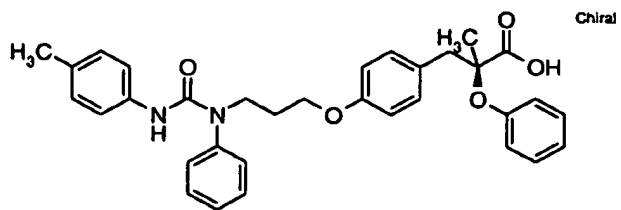
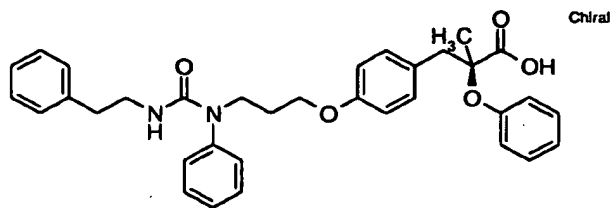


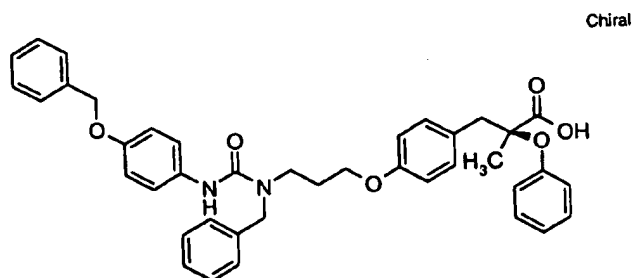
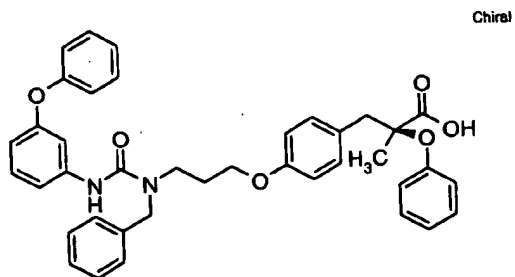
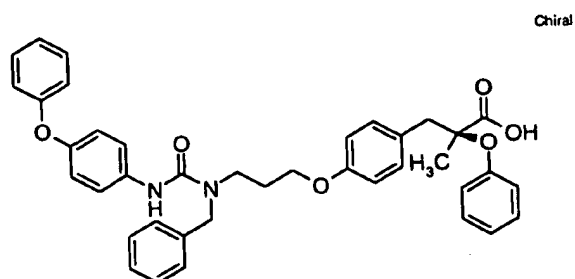
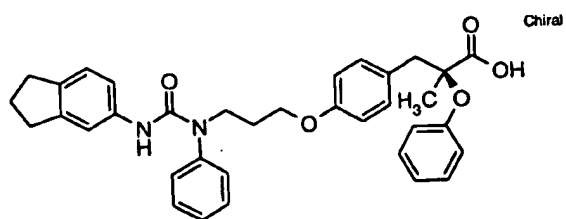
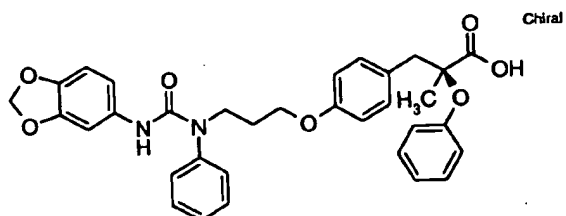
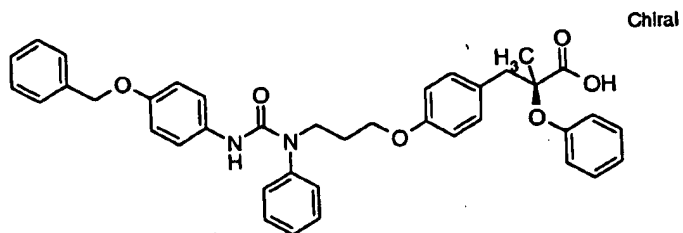












Docket No. X-15070

